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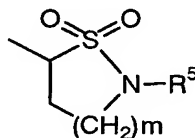
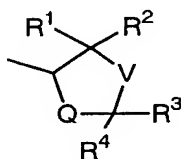


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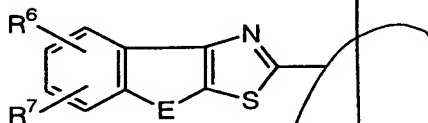
Z<sup>1</sup> is optionally substituted arylene, optionally substituted heteroarylene, optionally substituted non-aromatic heterocycle-diyl, or optionally substituted cycloalkyl-diyl;



oxygen atom or a sulfur atom; R<sup>3</sup> and R<sup>4</sup> are both hydrogen atoms or taken together may form an oxygen atom or a sulfur atom; R<sup>5</sup> is a hydrogen atom or lower alkyl; Q and V are each independently -O-, -S-, -NR<sup>B</sup>- (wherein R<sup>B</sup> is a hydrogen atom or lower alkyl), or -CH<sub>2</sub>-; m is 1, 2, or 3;

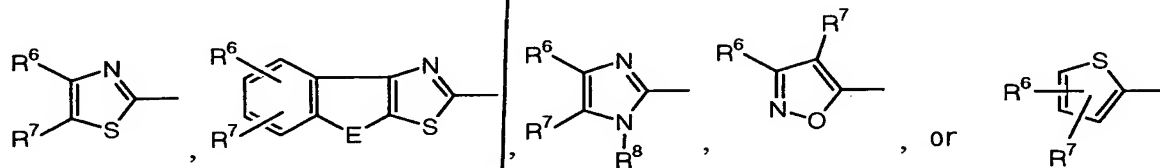
5 a broken line (---) represents the presence or absence of a bond,  
its prodrug, or their pharmaceutically acceptable salt, or solvate thereof.

2. A pharmaceutical composition exhibiting thrombopoietin agonism which contains a compound of claim 1, wherein X<sup>1</sup> is optionally substituted 5-member heteroaryl or a group represented by the formula:



10 wherein E is -(CH<sub>2</sub>)<sub>1-3</sub>-, -O-CH<sub>2</sub>-, or -S-CH<sub>2</sub>-; R<sup>6</sup> and R<sup>7</sup> are each independently a hydrogen atom, optionally substituted lower alkyl, carboxy, lower alkyloxycarbonyl, optionally substituted aminocarbonyl, optionally substituted thienyl, or optionally substituted phenyl.

15 3. A pharmaceutical composition exhibiting thrombopoietin agonism which contains a compound of claim 1, wherein X<sup>1</sup> is a group represented by the formula:

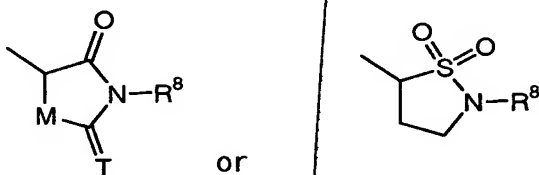


20 wherein E is -(CH<sub>2</sub>)<sub>1-3</sub>-, -O-CH<sub>2</sub>-, or -S-CH<sub>2</sub>-; R<sup>6</sup> and R<sup>7</sup> are each independently a hydrogen atom, optionally substituted lower alkyl, carboxy, lower alkyloxycarbonyl, optionally substituted aminocarbonyl, optionally substituted thienyl, or optionally substituted phenyl; R<sup>8</sup> is a hydrogen atom or lower alkyl.

4. A pharmaceutical composition of any one of claims 1 to 3, wherein Y<sup>1</sup> is -NHCO-, -CONH-, -NHCH<sub>2</sub>-, or -NHSO<sub>2</sub>-.

5. A pharmaceutical composition of any one of claims 1 to 4, wherein Z<sup>1</sup> is 1,4-phenylene.

6. A pharmaceutical composition of any one of claims 1 to 6, wherein A<sup>1</sup> is a ring represented by the formula:



wherein R<sup>8</sup> is a hydrogen atom or lower alkyl; M is -S-, -O-, -N(R<sup>c</sup>)-, or -CH<sub>2</sub>- (wherein R<sup>c</sup> is a hydrogen atom or lower alkyl); T is an oxygen atom or a sulfur atom.

7. A pharmaceutical composition of any one of claims 1 to 6, wherein the broken line represents the presence of a bond.

8. A pharmaceutical composition of any one of claims 1 to 7, which is for treating or preventing hemopathy.

9. A pharmaceutical composition of any one of claims 1 to 7, which is a platelet production modifier.

10. Use of a compound of any one of claims 1 to 7 for preparation of a pharmaceutical composition for treating hemopathy.

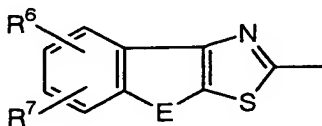
11. A method for treating hemopathy of a mammal, including a human, which comprises administration to said mammal of a compound of any one of claims 1 to 7 in a pharmaceutically effective amount.

12. A compound represented by the formula (II)



wherein X<sup>2</sup> is optionally substituted 5-member heteroaryl or a group

represented by the formula:



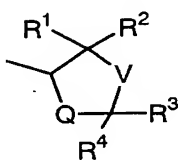
wherein E is  $-(CH_2)_{1-3}-$ ,  $-O-CH_2-$ , or  $-S-CH_2-$ ;  $R^6$  and  $R^7$  are each independently a hydrogen atom, optionally substituted lower alkyl, carboxy, lower alkyloxycarbonyl, optionally substituted aminocarbonyl, optionally substituted thienyl, or optionally substituted phenyl;

$Y^2$  is  $-NR^GCO-(CH_2)_{0-2}-$ ,  $-NR^GCO-(CH_2)_{0-2}-W-$ ,  $-NR^GCO-CH=CH-$ ,  $-W-(CH_2)_{1-5}-NR^GCO-(CH_2)_{0-2}-$ ,  $-W-(CH_2)_{1-5}-CONR^G-(CH_2)_{0-2}-$ ,  $-CONR^G-(CH_2)_{0-2}-$ ,  $-(CH_2)_{0-5}-NR^G-SO_2-(CH_2)_{0-5}-$ ,  $-(CH_2)_{0-5}-SO_2-NR^G-(CH_2)_{0-5}-$ ,  $-NR^G-(CH_2)_{0-2}-$ ,  $-NR^G-CO-NR^G-$ ,  $-NR^G-CS-NR^G-$ ,  $-N=C(-SR^G)-NR^G-$ ,  $-NR^GCSNR^GCO-$ ,  $-N=C(-SR^G)-NR^GCO-$ ,  $-NR^G-(CH_2)_{1-2}-NR^GCO-$ ,  $-NR^GCONR^GNR^FCO-$ , or  $-N=C(-NR^GR^G)-NR^GCO-$ ,

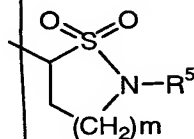
wherein  $R^G$  is each independently a hydrogen atom or optionally substituted lower alkyl,  $R^F$  is a hydrogen atom or optionally substituted aryl, W is an oxygen atom or a sulfur atom;

$Z^2$  is optionally substituted phenylene, optionally substituted 2,5-pyridine-diyl, optionally substituted 2,5-thiophene-diyl, or optionally substituted 2,5-furan-diyl;

$A^2$  is a ring represented by the formula:



or



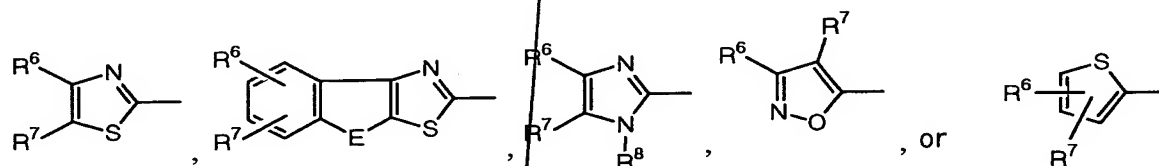
wherein  $R^1$  and  $R^2$  are both hydrogen atoms or taken together may form an oxygen atom or a sulfur atom;  $R^3$  and  $R^4$  are both hydrogen atoms or taken together may form an oxygen atom or a sulfur atom;  $R^5$  is a hydrogen atom or

lower alkyl; Q and V are each independently -O-, -S-, -NR<sup>B</sup>- (wherein R<sup>B</sup> is a hydrogen atom or lower alkyl), or -CH<sub>2</sub>-; m is 1, 2, or 3;

a broken line (---) represents the presence or absence of a bond,  
provided that X<sup>2</sup> is not oxazole,

its prodrug, or their pharmaceutically acceptable salt, or solvate thereof.

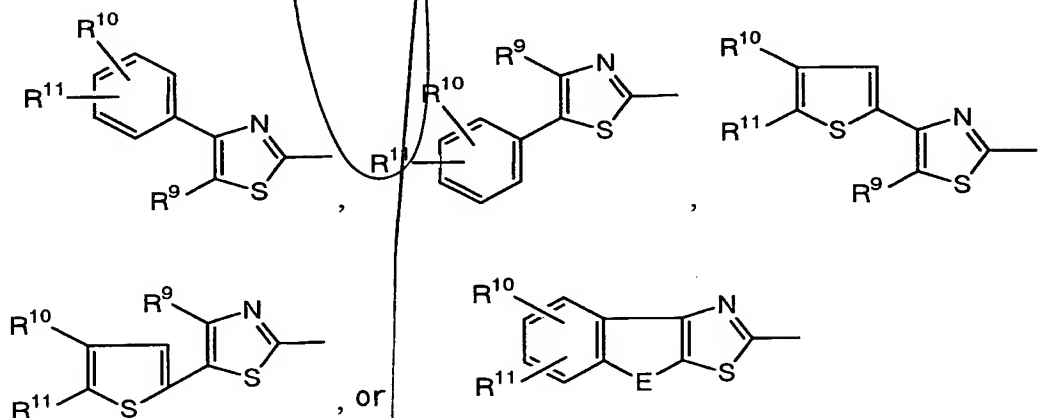
13. A compound of claim 12, wherein X<sup>2</sup> is a group represented by the formula:



wherein E is -(CH<sub>2</sub>)<sub>1-3</sub>-, -O-CH<sub>2</sub>-, or -S-CH<sub>2</sub>-; R<sup>6</sup> and R<sup>7</sup> are each independently a hydrogen atom, optionally substituted lower alkyl, carboxy, lower alkyloxycarbonyl, optionally substituted aminocarbonyl, optionally substituted thienyl, or optionally substituted phenyl; R<sup>8</sup> is a hydrogen atom or lower alkyl,

its prodrug, or their pharmaceutically acceptable salt, or solvate thereof.

14. A compound of claim 12, wherein X<sup>2</sup> is a group represented by the formula:



wherein E is as defined in claim 12;

R<sup>9</sup> is a hydrogen atom, optionally substituted lower alkyl, carboxy, lower

alkyloxycarbonyl, or optionally substituted aminocarbonyl;

R<sup>10</sup> and R<sup>11</sup> are each independently a hydrogen atom, halogen, carboxy, lower alkyloxycarbonyl, optionally substituted aminocarbonyl, nitro, or optionally substituted amino,

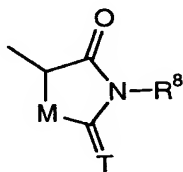
5 its prodrug, or their pharmaceutically acceptable salt, or solvate thereof.

15. A compound of any one of claims 12 to 14, wherein Y<sup>2</sup> is -NHCO-, -CONH-, -NHCH<sub>2</sub>-, or -NHSO<sub>2</sub>-,

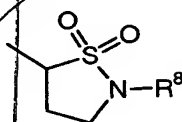
its prodrug, or their pharmaceutically acceptable salt, or solvate thereof.

10 16. A compound of any one of claims 12 to 15, wherein Z<sup>2</sup> is 1,4-phenylene, its prodrug, or their pharmaceutically acceptable salt, or solvate thereof.

17. A compound of any one of claims 12 to 16, wherein A<sup>2</sup> is a ring represented by the formula:



or



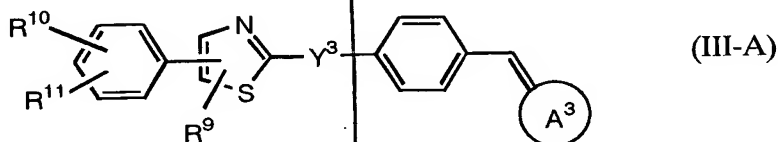
15 wherein R<sup>8</sup> is a hydrogen atom or lower alkyl; M is -S-, -O-, -N(R<sup>c</sup>)-, or -CH<sub>2</sub>- (wherein R<sup>c</sup> is a hydrogen atom or lower alkyl); T is an oxygen atom or a sulfur atom,

its prodrug, or their pharmaceutically acceptable salt, or solvate thereof.

18. A compound of any one of claims 12 to 17, wherein the broken line represents the presence of a bond,

20 its prodrug, or their pharmaceutically acceptable salt, or solvate thereof.

19. A compound represented by the formula III-A:



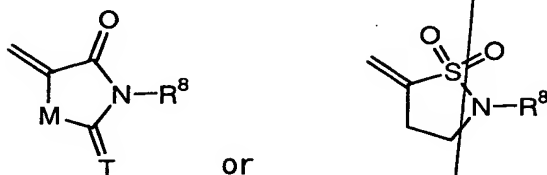
wherein, R<sup>9</sup> is a hydrogen atom, optionally substituted lower alkyl, carboxy,

lower alkyloxycarbonyl, or optionally substituted aminocarbonyl;

R<sup>10</sup> and R<sup>11</sup> are each independently a hydrogen atom, halogen, carboxy, lower alkyloxycarbonyl, optionally substituted aminocarbonyl, nitro, or optionally substituted amino;

5 Y<sup>3</sup> is -NHCO- or -CONH-;

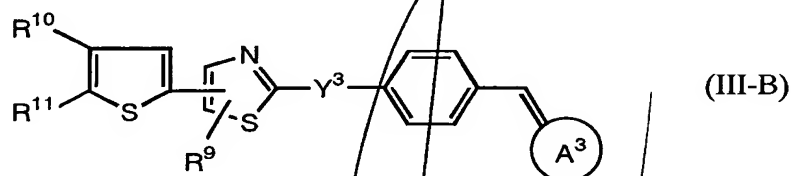
A<sup>3</sup> is a ring represented by the formula:



wherein R<sup>8</sup> is a hydrogen atom or lower alkyl; M is -S-, -O-, -N(R<sup>c</sup>)-, or -CH<sub>2</sub>- (wherein R<sup>c</sup> is a hydrogen atom or lower alkyl); T is an oxygen atom or a sulfur atom,

its prodrug, or their pharmaceutically acceptable salt, or solvate thereof.

20. A compound represented by the formula III-B:



wherein R<sup>9</sup>, R<sup>10</sup>, R<sup>11</sup>, Y<sup>3</sup>, and A<sup>3</sup> ring are as defined in claim 19,

its prodrug, or their pharmaceutically acceptable salt, or solvate thereof.

21. A pharmaceutical composition containing a compound of any one of claims 12 to 20 as an active ingredient.

22. A pharmaceutical composition which contains as an active ingredient a compound of any one of claims 12 to 20 for exhibiting thrombopoietin agonism.

23. An agent for treating or preventing hemopathy which contains as the active ingredient a compound of any one of claims 12 to 20.

24. A pharmaceutical composition containing as the active ingredient a

compound of any one of claims 12 to 20, which is a platelet production modifier.

25. Use of a compound of any one of claims 12 to 20 for preparation of a pharmaceutical composition for treating hemopathy.

5 26. A method for treating hemopathy of a mammal, including a human, which comprises administration to said mammal of a compound of any one of claims 12 to 20 in a pharmaceutically effective amount.



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